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REQUEST FOR FILING APPLICATION

Under Rule 53(a), (b)(1) & (d)(1)

(No Filing Fee or Oath/Declaration)

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Use for Design or Utility Applications

PATENT
APPLICATION

RULE 53(d) NO DECLARATION

Honorable Commissioner of
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Washington, DC 20231

Atty. Dkt.

240606

M#

D879PR

Client Ref

Date:

October 30, 1997

Sir:

1. This is a Request for filing a new Patent Application (☐ Design ☒ Utility) entitled:2. (Complete) Title: **DIELECTRIC, RADIATION-CURABLE COATING COMPOSITIONS AND METAL CONDUCTORS COATED WITH SUCH COATING**without a filing fee or Oath/Declaration but for which is enclosed the following:3. ☒ Abstract 1 page(s).4. 34 Pages of Specification (only spec. and claims); 5. ☐ Specification in non-English language6. 23 Numbered claim(s); and7. ☒ 6 sheet(s) per set: ☐ 1 set informal; 8. ☒ formal of size: ☒ A4 ☐ 11"
Drawings:9. **DOMESTIC/INTERNATIONAL** priority is claimed under 35 USC 119(e)/120/365(c) based on the following provisional, nonprovisional and/or PCT international application(s):

Application No.	Filing Date	Application No.	Filing Date
(1) 60/029,780	10/31/96	(2)	
(3)		(4)	
(5)		(6)	

10. **FOREIGN** priority is claimed under 35 USC 119(a)-(d)/365(b) based on filing in

Application No.	Filing Date	Application No.	Filing Date
(1)		(2)	
(3)		(4)	
(5)		(6)	

11. (No.) Certified copy (copies): ☐ attached; ☐ previously filed (date)
in U.S. Application No. / filed on 12. ☐ This is a reissue of Patent No. 13. ☐ **Amend the specification** by inserting before the first line - This is a ☐ Continuation-in-Part
☐ Divisional ☐ Continuation ☐ Substitute Application (MPEP 201.09) of:13(a) ☐ National Appln. No. / filed . - - (M#)13(b) ☐ International Appln. No. PCT/ filed which designated the U.S. - -14. ☐ See top of first page re continuing appln (X box only if info is there)14(a) extension to date: ☐ concurrently filed ☒ not needed ☐ previously filed

15. ☐: Prior application is assigned to

by Assignment recorded _____ Reel _____ Frame _____

16. ☐ Attached:

17. This application is made by the following named inventor(s) **(Double check instructions for accuracy):**

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18. NOTE: FOR ADDITIONAL INVENTORS, check box ☐
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APPLICATION UNDER UNITED STATES PATENT LAWS

Invention: DIELECTRIC, RADIATION-CURABLE COATING COMPOSITIONS AND METAL CONDUCTORS COATED WITH SUCH COATING

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This is a:

- ☐ Provisional Application
- ☒ Regular Utility Application
- ☐ Continuing Application
- ☐ PCT National Phase Application
- ☐ Design Application
- ☐ Reissue Application
- ☐ Plant Application
- ☐ Substitute Specification

Sub. Spec Filed _____
in App. No. ____ / _____

SPECIFICATION

**DIELECTRIC, RADIATION-CURABLE COATING COMPOSITIONS AND
METAL CONDUCTORS COATED WITH SUCH COATING**

1. Field of the Invention

5 This invention relates to radiation-curable coating compositions for coating metal conductors, and in particular, to coating compositions which provide good insulating properties in extreme conditions.

10 **2. Description of Related Art**

 Metal conductors are in general coated with a dielectric coating for insulating the conductor. Such coatings require good insulating properties in a variety of environments. It is particularly important
15 for such coatings to provide good insulating properties under extreme conditions such as in transformer coils found in power distribution transformers. U.S. patent 4,481,258 issued to Sattler et al. discloses the use of paper as insulating material. Although Sattler
20 proposes a coating be used as insulating material, it fails to disclose a coating sufficient to meet the requirements of an insulator in extreme conditions. As a result, paper insulation materials are still being used in the manufacture of transformer coils. The
25 coatings proposed in Sattler are certain UV-curable materials comprising acrylate-ester adducts, acrylate urethane adducts and acrylate functional diluents. These coatings require both UV cure, and an additional

post-cure at a temperature of 130°C for 4-17 hr. The use in transformer coils of the coatings and processes disclosed in Sattler is unattractive; in particular because of the post-cure required.

5 Effective insulating coatings for high power transformer coils should exhibit the desirable properties described below.

 As the metal is coated and thereafter is bent in a required form, the cured coating should be
10 flexible so that it can withstand bending of the coated conductor as it is wound into a coil.

 The cured coating should be able to withstand immersion in oil for 28 days at 150°C as described in US-A-4481258.

15 The cured coating should remain adherent at elevated temperature that is encountered when the transformer is under load.

 The cured coating should have a dielectric constant smaller than 5% at 60 Hz (24°C).

20 The cured coating should have a dielectric dissipation factor smaller than 0.05 at 24°C before and after hot oil exposure and smaller than 0.2 at 150°C, both at 60 Hz.

 It is an object of the present invention to
25 provide dielectric radiation-curable coating compositions for metal conductors which have the properties identified above.

It is further an object of the present invention to provide metal conductors which are coated with UV-curable coating composition having the properties identified above.

5 It is further an object of the present invention to disclose a method of manufacture of metal conductors which are coated with UV-curable coating composition having the properties identified above.

10 **3. Summary of the Invention**

The Invention relates to a metal conductor with a cured coating of about 2.5 μm to about 500 μm and preferably about 10 μm to about 50 μm thickness which cured coating has a dielectric dissipation factor
15 (60Hz, 24°C) of lower than about 0.05 and is a radiation-cured coating formulated from components comprising:

- a) an acrylate functional urethane oligomer having a hydrocarbon backbone
- 20 b) one or more mono- or polyfunctional diluents, and optionally,
- c) one or more light sensitive radical generating compounds.

Furthermore, the invention relates to a
25 radiation-curable coating composition for coating a metal conductor comprising:

- a) an acrylate functional urethane oligomer having a hydrocarbon backbone

b) one or more mono- or polyfunctional diluents; and
optionally

c) one or more light sensitive radical generating
compounds,

5 which coating, when cured with radiation, has a
dielectric dissipation factor at 60 Hz at 24°C of lower
than about 0.05, a dissipation factor at 60 Hz at 150°C
of lower than about 0.2, and an elongation at 25°C of a
25 μm thin coating of greater than about 50%.

10 The invention also relates to a method of
making a metal conductor with a cured coating of about
2.5 μm to about 500 μm and preferably about 10 μm to
about 50 μm thickness which cured coating has a
dielectric dissipation factor (60Hz, 24°C) of lower than
15 about 0.05 and is a radiation-cured coating formulated
from components comprising:

a) an acrylate functional urethane oligomer having a
hydrocarbon backbone

b) one or more mono- or polyfunctional diluents, and
20 optionally,

c) one or more light sensitive radical generating
compounds.

4. Brief Description of the Drawings

25 Figure 1 provides a graphical illustration of
temperature dependence of electrical dissipation factor
of the radiation-curable coating and the photoinitiator

concentration effect on the electrical dissipation factor for Example XI.

Figure 2 provides a graphical illustration of the phthalocyanine pigment effect on electrical dissipation factor for Example XII.

Figure 3a and 3b provide the elastic modulus (E'), viscosity modulus (E'') and tan delta peak (E''/E') as measured by conventional DMA methods for coating Examples X and XI of the present invention, respectively.

Figures 4a and 4b provide the average field strength at Dielectric breakdown for different thicknesses of coating Example XI of the present invention as measured with the applied potential being held constant and the applied potential increasing at the rate of 500V/second, respectively.

5. Detailed Description of the Invention

The cured coating layer on the metal conductor has outstanding insulating properties, both at low and high temperature. The insulating cured coating layer appears to have a low dielectric constant, e.g. lower than about 5 (60Hz, 24°C) and a good dielectric breakdown value. Furthermore, the cured coating is flexible as to allow bending of the metal conductor.

The metal conductor preferably is an iron,

not cause damage to the cured coating. Thus, the cured coating measured at 25 μm thickness, preferably has an elongation of at least about 50%. In particular, the coating has at least one T_g of below 20°C (as measured by the peak of the $\tan \delta$ curve in a Dynamic Mechanical Analysis (DMA) at 1 Hz.

As the coated metal conductor can be used in a hot oil environment, most preferably the cured coating has a dissipation factor at 60 Hz at 150°C of lower than about 0.2. Furthermore, the cured coating preferably has an electrical dissipation factor at 60 Hz at 24°C before and after a hot oil aging test of lower than about 0.05.

The cured coating of the present invention exhibits its insulating properties even when formed as a very thin film. The cured coating has a thickness of about 2.5 μm to about 50 μm , preferably between about 10 μm to about 100 and more preferably between about 10 μm to about 50 μm .

The first component of the radiation-curable coating is a hydrocarbon oligomer end-capped with acrylate functional urethane or acrylated acrylic groups. A preferred first component is an acrylate functional urethane oligomer (a) having a hydrocarbon backbone. The word backbone is used to denote the oligomer or polymer to which the acrylate urethane groups are attached. This acrylate functional urethane

oligomer preferably is used in an amount of about 20-80 wt.% with respect to the total coating composition. More preferably, the amount is about 30-65 wt.%.

5 The oligomer (a) utilized in the present invention preferably is the reaction product of (i) a hydrocarbon compound with groups reactive with an isocyanate; (ii) a polyisocyanate; and (iii) an hydroxy functional endcapping monomer.

10 The hydrocarbon compound with groups reactive with an isocyanate (i) is provided by a linear or branched hydrocarbon containing a plurality of said reactive end groups, and providing a hydrocarbon backbone to the oligomer. The isocyanate reactive groups may be thiol, amine or hydroxy. Particularly
15 preferred are hydroxy groups. Because of the amine and thiol groups, the urethane oligomer may comprise low concentrations of urea or thio-urea groups, for example below about 5 wt.%. The hydrocarbon portion is from about 200 to about 5,000 molecular weight and
20 preferably from about 400 to about 4,000 molecular weight. Molecular weight in this case is determined by gel permeation chromatography (GPC), using a methylene chloride solvent, as measured against polystyrene molecular weight standards. By "hydrocarbon" is meant
25 a non-aromatic compound containing a majority of methylene groups ($-\text{CH}_2-$) and which may contain internal unsaturation and/or pendent unsaturation. Fully saturated (i.e., hydrogenated) hydrocarbons are preferred because the electric dissipation factor of

the cured coating increases as the degree of unsaturation increases. Suitable hydrocarbon polyols include hydroxyl-terminated, fully or partially hydrogenated 1,2-polybutadiene; 1,4- and 1,2-polybutadiene copolymers; 1,2-polybutadiene polyol hydrogenated to an iodine number of from 9 to 21; fully or partially hydrogenated polyisobutylene polyol; mixtures thereof, and the like. Preferably, the hydrocarbon polyol is substantially fully hydrogenated, and thus a preferred polyol is hydrogenated 1,2-polybutadiene, and hydrogenated 1,4-,1,2-polybutadiene copolymers with about 50 wt.% to about 80 wt.% 1,4-butadiene and about 20 wt.% to about 50 wt.% 1,2-butadiene copolymerized monomers. Suitable hydrocarbon polyamines or polythiols include the above described polyols with thiol or amino groups instead of the hydroxy groups.

The polyisocyanate component (ii) is aromatic or non-aromatic, and preferably is non-aromatic. A suitable aromatic polyisocyanate is toluene diisocyanate. Non-aromatic polyisocyanates of from 4 to 20 carbon atoms may be employed. Suitable saturated aliphatic polyisocyanates include isophorone diisocyanate; dicyclohexylmethane-4,4'-diisocyanate; 1,4-tetramethylene diisocyanate; 1,5-pentamethylene diisocyanate; 1,7-heptamethylene diisocyanate; 1,8-octamethylene diisocyanate; 1,9-nonamethylene diisocyanate; 1,10-decamethylene diisocyanate; 2,2,4-trimethyl-1,5-pentamethylene diisocyanate; 2,2'-

dimethyl-1,5-pentamethylene diisocyanate; 3-methoxy-
1,6-hexamethylene diisocyanate; 3-butoxy-1,6-
hexamethylene diisocyanate; omega,omega'-dipropylether
diisocyanate; 1,4-cyclohexyl diisocyanate; 1,3-
5 cyclohexyl diisocyanate; trimethylhexamethylene
diisocyanate; and mixtures thereof. Isophorone
diisocyanate is the preferred aliphatic polyisocyanate.

The reaction rate between a hydroxyl-
terminated hydrocarbon and the diisocyanate may be
10 increased by use of a catalyst in the amount of about
100 ppm to about 200 ppm. Suitable catalyst include
dibutyl tin dilaurate, dibutyl tin oxide, dibutyl tin
di-2-hexanoate, stannous oleate, stannous octoate, lead
octoate, ferrous acetoacetate, and amines such as
15 triethylamine, diethylmethanamine, triethylenediamine,
dimethyl-ethylamine, morpholine, N-ethyl morpholine,
piperazine, N,N-dimethyl benzylamine, N,N-dimethyl
laurylamine, and mixtures thereof. A preferred
catalyst is dibutyl tin dilaurate.

20 The endcapping monomer (iii) is a hydroxyl-
terminated aliphatic acrylate or methacrylate,
preferably an alkoxyated (meth)acrylate wherein 1-10
molecules of ethylene, propylene or butylene oxide are
reacted with acrylic or methacrylic acid.

25 Suitable hydroxyl-terminated monoacrylates
which may be used as the endcapping monomer include
hydroxyethyl acrylate, hydroxyethyl methacrylate,
hydroxypropyl acrylate and hydroxypropyl methacrylate.
Hydroxyethyl acrylate is preferred because it imparts a

1
faster cure rate to the polyurethane oligomer. The
molar ratio of the hydrocarbon compound, diisocyanates
and endcapping monomer is preferably approximately
1:2:2.

5 The second component (b) is constituted by
one or more mono- or polyfunctional diluents.
Preferably, the diluents are acrylate or methacrylate
functional. However, minor amounts of other types of
monomers can be used as well. The amount of component
10 (b) preferably is about 20 wt.% to about 80 wt.% of the
total coating composition, more preferred about 20 wt.
% to about 70 wt.%. Particularly preferred is the use
of about 10 wt.% to about 50 wt.% of monofunctional
diluent(s), and about 5 wt. % to about 40 wt.% of
15 polyfunctional diluent(s).

 The second component (b) of the composition
is a monomer which preferably comprises a
monofunctional alkyl acrylate or methacrylate-based
monomer. The alkyl portion of the monomer has between
20 6 and 18 carbon atoms, and preferably between 8 and 15
carbon atoms, and therefore is hydrocarbon in
character. This monomer may be either straight chain,
branched or cyclic. This component comprises from
about 5 wt. % to about 50 wt. % of the composition,
25 based upon the total weight of the coating composition.
Preferably, it comprises from about 5 wt. % to about 50
wt. %, and more preferably from about 10 wt. % to about
40 wt. % of the composition.

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The monomer, component (b) is selected to be one that is compatible with the oligomer discussed above. Suitable examples of C₆ to C₁₈ alkyl acrylate or methacrylate-based monomers include hexyl acrylate; 5 hexyl methacrylate; cyclohexylacrylate; cyclohexylmethacrylate; 2-ethylhexyl acrylate; 2-ethylhexyl methacrylate; isooctyl acrylate; isooctyl methacrylate; octyl acrylate; octyl methacrylate; decyl acrylate; decyl methacrylate; isodecyl acrylate; isodecyl 10 methacrylate; isobornylacrylate; isobornylmethacrylate; lauryl acrylate; lauryl methacrylate; stearyl acrylate; stearyl methacrylate.

The second component, component (b), further comprises preferably an alkylacrylate polyfunctional 15 diluent (or monomer) in an amount of about 5 wt. % to about 50 wt.%, preferably about 5 wt. % to about 40 wt.%. Suitable examples of these polyfunctional monomers are C₄-C₁₅ hydrocarbon diol acrylates; C₄-C₁₅ hydrocarbon diol methacrylates; and mixtures of the 20 above. The term hydrocarbon includes cycloalkylgroups. Other suitable polyfunctional acrylates are (alkoxylated) polyolpolyacrylates. Examples of suitable polyfunctional monomers include butanediol dimethacrylate, butanediol diacrylate, propanediol 25 dimethacrylate, propanediol diacrylate, pentanediol dimethacrylate, pentanediol diacrylate, hexanediol dimethacrylate, hexanediol diacrylate, neopentylglycol dimethacrylate, neopentylglycol diacrylate, trimethylolpropane triacrylate, trimethylolpropane

trimethacrylate, polypropylene glycol diacrylate, polypropylene glycol dimethacrylate, cyclohexanedi-methanoldiacrylate or -methacrylate and, tricyclodecane dimethanol di(meth)acrylate.

5 Preferred alkyl acrylate monomers include isobornyl acrylate, 2-ethylhexylacrylate, isooctylacrylate, cyclohexylacrylate, hexanedioldiacrylate, tricyclodecanedimethanol-diacrylate.

10 Other diluents may be used in amounts of preferably less than about 10 wt.%. Examples of these diluents are N-vinyl functional or vinyl ether functional compounds with a molecular weight lower than about 500. Examples of these diluents are N-vinyl
15 caprolactam, butyl-vinyl ether, triethyleneglycoldi-vinyl ether, butanediol-divinyl ether and the like.

The diluents preferably are used in a quantity sufficient to adjust the total coating composition to a viscosity of lower than about 2000
20 mPa.s, preferably lower than about 800 mPa.s at 25°C, measured with a Couette apparatus (cup-and-bob viscometer at a frequency 100 rpm).

The coating composition of the present invention preferably does not comprise substantial
25 amounts of monomers with relatively strong dipole moments such as N-vinylpyrrolidone, phenoxyethylacrylate, polyoxyalkylene-alkylphenolacrylate and the like. The coating

composition furthermore, preferably does not comprise, in substantial amounts, those monomers for which dipoles can be easily included, such as aromatic groups containing acrylates such as phenylacrylates. The person skilled in the art can easily determine the amount allowed in the composition by measuring the dissipation factor.

The coating composition is radiation-curable, and can be cured with electron beam irradiation or with light with a wavelength between about 200 nm and about 700 nm. In the latter case, the composition comprises a light sensitive radical generating compound or mixture of compounds which act as photoinitiators.

The photoinitiator, when used in a small but effective amount to promote radiation cure, must provide reasonable cure speed without causing premature gelation of the composition.

Suitable photoinitiators include the following: hydroxycyclohexylphenyl ketone; hydroxymethylphenylpropanone; dimethoxyphenylacetophenone; 2-methyl-1, [4-(methyl thio)phenyl]-2-morpholino-propanone-1; 1-(4-isopropylphenyl)-2-hydroxy-2-methylpropan-1-one; 1-(4-dodecylphenyl)-2-hydroxy-2-methylpropan-1-one; 4-(2-hydroxyethoxy)phenyl- (2-hydroxy-2-propyl)ketone; diethoxyacetophenone; 2,2-di-sec-butoxyacetophenone; diethoxy-phenyl acetophenone; and mixtures of these.

The photoinitiator, if used, preferably comprises from about 1.0 percent to about 10.0 percent

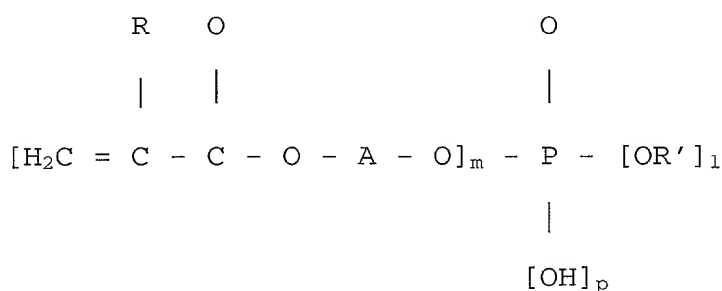
by weight of the composition, based upon the total composition. Preferably, the amount of photoinitiator is from about 2.0 percent to about 7.0 percent by weight. The photoinitiator should be chosen such that a cure speed, as measured in a dose versus modulus curve, of less than about 2.0 J/cm², and preferably less than about 1.0 J/cm², is required, when the photoinitiator is used in the designated amount.

FIG. 1 shows the effect on dielectric dissipation factor of cured coatings when varying amounts of photoinitiator are added to pre-cured coating compositions for Example X (3 wt.% photoinitiator) and Example XI (6 wt.% photoinitiator).

The composition preferably also contains an adhesion promoter. The adhesion promoter is preferably a compound having a groups participating in the radical curing reaction, and a group that adheres to the metal conductor. The group that participates in the curing reaction can be preferably, vinyl, (meth) acrylate or thiol. The group that adheres to the metal conductor preferably is hydroxy, acid, zirconate, titanate or silane. The acid may be for example carboxylic phosphoric or sulphonic. Most preferred is a (meth)acrylate functionalized carboxylic acid or phosphoric acid. Some examples of suitable adhesion promoters include, but are not limited to, hydroxyethyl(meth)acrylate, hydroxypropyl(meth)acrylate, di- or trialkoxy zirconates or titanates, vinyl trimethoxysilane, mercaptopropyltrimethoxy

silane, acrylic acid, methacrylic acid, β -carboxyethyl
acrylate, EBERCYL 170 and EBERCYL 169. The EBERCYL
products are acrylate ester derivatives, available from
Radcure Specialties in Atlanta, Georgia, and are
5 phosphoric acid based adhesion promoters.

Mono or diester or phosphoric acid having the
following formula are also suitable adhesion promoters:



where

$$m + 1 + p = 3$$

$$R = H \text{ or } CH_3$$

$$A = C_nH_{2n}, \text{ and } 2 \leq n \leq 6$$

$$R' = C_1 \text{ to } C_{14} \text{ alkyl, aryl, alkaryl, or alkyleneoxy.}$$

Representative of the various species of
organo-phosphate esters having the above formula
include, but are not limited to,

(1) methylmethacryloyloxyethyl phosphate, where

$$\begin{array}{l}
 (R = CH_3; A = -C_2H_4-; R' = CH_3, m = 1 \text{ and} \\
 p = 1);
 \end{array}$$

(2) ethyl methacryloyloxyethyl phosphate, where

$$(R = CH_3; A = -C_2H_4-; R' = C_2H_5); m = 1 \text{ and}$$

p = 1);

(3) propylacryloyloxyethyl phosphate, where (R = H;

A = -C₂H₄-; R' = C₃H₇; m = 1 and p = 1);

(4) methyl acryloyloxyethylphosphate, where (R = H,

5 A = -C₂H₄-; R' = CH₃, m = 1 and p = 1);

(5) ethylacryloyloxyethylphosphate, where (R = H;

A = -C₂H₄-; m = 1 and p = 1; R' = C₂H₅);

(6) propylmethacryloyloxy-ethylphosphate, where

(R = CH₃; A = -C₂H₄-; R' = C₃H₇; m = 1 and

10 p = 1);

(7) bis(methacryloyloxyethyl) phosphate, where (R = CH₃;

A = -C₂H₄-; m = 2; l = 0; p = 1); and

(8) bis(acryloyloxyethyl)phosphate, where (R = H;

A = -C₂H₄-; m = 2; l = 0; p = 1).

15 The adhesion promoter helps the coating composition adhere to the metal conductor. The adhesion promoter may be used in an amount in the range of about 0.2 wt.% to about 5 wt.% of the composition. Care should be exercised, that the amount of adhesion
20 promotor is not so large that insulating properties are decreased below acceptable level.

It is an unexpected advantage of the coating composition of the present invention, that an adhesion promoter can be used in effective amounts while
25 maintaining very good insulating properties for the cured coating.

In addition to the above components, the composition may also contain other components that are known to those skilled in the art including

stabilizers, surfactants, plasticizers, chain transfer agents and the like.

In addition, it may be useful to use a small amount of pigment or dye to color the cured coating.

5 This allows simple visual control of the coated metal conductor. This is in particular useful, in case the metal conductor is only partly coated. Suitable pigments or dyes are for example copper phthalocyanine blue, crystal violet lactone (blue), crystal malachite
10 green, sheet fed rubine (red). The amount of pigment, if used, will in general be about 0.2 wt. % to about 5 wt.% relative to the coating composition.

FIG. 2 shows the dielectric dissipation factor for sample coatings, Examples XI and XIII, and
15 the absence of any negative effect resulting from the addition of phthalocyanine pigment (0.5 wt.%) to the pre-cured coating composition.

The coating composition may be applied on the metal conductor using known coating method, such as
20 spraying, vacuum coating, dipping and doctoring. The coating composition may be applied under a nitrogen atmosphere to preclude oxygen inhibition; however, this is not strictly necessary. If, for example, a relatively large amount of photoinitiator is used in
25 the composition, the cure of the surface of the film is adequate as well.

The invention will be further demonstrated by the following, non limiting examples.

Examples

Preparation of acrylate functional oligomer A.

5 Isophorone diisocyanate (IPDI 429 g) was
dissolved in laurylacrylate (420 g) with 1 g BHT
(butylated hydroxy toluene) 0.7 g phenolthiazine and 2
g dibutyltindilaurate. To this mixture, 224 g of
hydroxyethylacrylate (HEA) was slowly added, and the
10 temperature was kept below 35°C. To the acrylate-
isocyanate adduct, 2318 g of a hydrocarbon diol was
added (Nisso PB 2000) and it was allowed to react.
About 105 g of laurylacrylate was added and the final
NCO content was determined to be below 0.1 %. The
15 resulting oligomer A had a theoretical molecular weight
of 3089 and was a clear solution of 85% oligomer in 15%
laurylacrylate.

Preparation of acrylate functional oligomer B

20 In an analogous way as the preparation of
oligomer A, an oligomer was made from 400 g IPDI, 139 g
HEA, 2876 g Nisso PB 2000 and 380 g laurylacrylate.
The theoretical molecular weight of the resulting
oligomer B is 5733.

Preparation of acrylate functional oligomer C

25 In an analogous way as the preparation of
oligomer A, oligomer C was prepared from 81 g IPDI, 42

g HEA, 430 g of Nisso PB 2000 and 140 g isobornylacrylate. The theoretical molecular weight of the resulting oligomer C is 3093.

5 Examples I-VII

 Coating compositions were prepared with the oligomer in 15% diluent mixtures using oligomers A-C, with further diluents and a photoinitiator as shown in Table 1. The coating compositions were applied on an aluminum plate and cured with 2 J/cm² light of a fusion D bulb. For measuring the dissipation factor 150 μm thick films were cast on a glass plate and cured with 2 J/cm; the dissipation factor was measured at 24°C and 150°C at 60 Hz with standard equipment with stainless steel electrodes. Results are shown in Table 1.

TABLE 1

Component	I	II	III	IV	V	VI	VII
oligomer A	51.8		57		57		
oligomer B		51.8		57		57	57
oligomer C							
laurylacrylate	36.4	36.4	30	30	30	30	30
isobornylacrylate							
PHOTOMER 3016 ¹⁾	9.1	9.1	10	10	10	10	10
SA 1002 ²⁾							
SR 349 ³⁾							
IRGACURE 500 ⁴⁾	2.7	2.7	3	3	3	3	3
Dissipation factor 60 Hz							
at 24°C	0.028	0.038	0.030	0.033	0.033	0.015	0.027
at 150°C	0.098	0.106	0.021	0.023	0.023	0.010	0.028
Dielectrical constant	< 3.0	< 3.0	< 3.0	< 3.0	< 3.0	< 3.0	< 3.0

¹⁾ PHOTOMER 3016 is: Bisphenol-A-diacrylate available from Henkel

²⁾ SA 1002 is: tricyclododecanedimethanol diacrylate available from Sartomer

³⁾ SR349 is: ethoxylated bis phenol-A-diacrylate available from Sartomer

⁴⁾ IRGACURE 500 is available from Ciba-Geigy

Examples VIII-XVIII

In an analogous way, further coating compositions were made and tested. The oligomer C was used in these examples.

5 Compositions and results are summarized in Tables 2 and 3. The coating compositions were spin coated on an aluminum panel, and cured with 1 J/cm, resulting in a 12.5 μm film; furthermore, coatings were cast on a glass plate and cured with 2 J/cm².

10

Test Procedures

Elastic modulus (E'), viscous modulus (E''), and tan delta (E''/E') were measured by conventional DMA methods. A free film specimen of cured material
15 was cut to size (about 35 mm length), measured for width and thickness and mounted. The environmental chamber containing the sample was brought to 80°C. The sample was stretched before temperature sweep was started. The temperature was lowered in prescribed
20 steps to the starting temperature. The temperature sweep was started and allowed to proceed upwards on the temperature scale until the material was beyond the glass transition range and well into the rubbery range. The DMA instrument (Rheometrics Solids Analyzer, RSA-II
25 equipped with a personal computer) produced a plot of the data on the computer screen. The temperature at which E' is 1,000 MPa and E' is 100 MPa was calculated from this plot, as well as the tan delta peak. The

minimum value of E' attained in the rubbery region was measured.

Cured coating samples were tested for flexibility and structural integrity under the strain required for a 180° bend of coated 0.0625" thick aluminum substrate over 0.25" mandrel at 25°C. Visual examination of the tested samples indicated the cured samples tested maintained structural integrity and did not delaminate from the aluminum substrate. Results of the 180° bend test are recorded for Examples VIII, IX, X, XI, XII, XIII in Table 2. Results for Examples XIV, XV, XVI, XVII and XVIII are shown in Table 3.

Samples of the coating compositions were tested for blocking by spin coating the pre-cured compositions on aluminum panels to form films. The compositions were cured at 1 J/cm², under air and testing at 60°C by placing weights on top of a 10 x 10 cm area of superimposed films. The samples were then placed in the oven for 3 days. No blocking was detected. The results shown in Tables 2 and 3.

Samples of 25μm thick cured coating compositions were tested for adhesion to a 1.59 mm thick aluminum panel which was bent 180° over a 6.4mm diameter mandrel and immersed in 150°C oil for 28 days. Adhesion of the 25μm thick cured coating samples was tested using crosshatch method well known in the art and described in detail in "Coating Technology

Handbook", by U. Zorll, published by Marcel Dekker, Inc., (1991) and incorporated herein by reference. The results are shown in Tables 2 and 3.

Dissipation factor of the cured coatings was tested by methods well known in the art. Results are recorded in Tables 2 and 3. Figures 1 and 2 provide additional dielectric dissipation factor recordings for cured coating composition Examples X, XI and XIII.

The temperature sensitivity of dissipation factors (%) for Examples IX, X, XI and XIII for temperatures ranging from 25°C through 185°C are shown in Table 4.

TABLE 2

Component	VIII	IX	X	XI	XII	XIII
Oligomer C	50	50	55.6	54.03	48.45	53.8
Isobornylacrylate			11.1	10.8		10.8
tricyclodecanedimethanol diacrylate	24.0	24.0			23.26	
2-ethylhexylacrylate		20.0				
hexanedioldiacrylate			26.7	25.93		25.8
cyclohexylacrylate	20.0					
isooctylacrylate					19.38	
DAROCURE 1173 (Ciba-Geigy)	3.0	3.0	3.3	6.0	6.0	5.9
EBECRYL 170 (Radcure)	3.0	3.0	3.3	3.24	2.91	3.2
phthalocyanine blue						0.5
Viscosity (mPa.s) at 25°C	700	500	785	730	450	730
Dissipation Factor at 25°C, 60Hz	0.02	0.029	0.019	0.024	~ 0.02	0.025
Dissipation Factor at 150°C, 60Hz	0.083	0.032	0.060	0.082	~ 0.05	0.053
stability under 150°C oil immersion	pass	pass	slight defect	pass	pass	pass
Blocking of coating at 60°C	nd	nd	nd	pass	pass	pass
Dielectric constant at 24°C, 60 Hz	< 2.7	< 2.7	< 2.7	< 2.7	< 2.7	< 2.7
Tg (°C)		-17	-11			
180° bend test	pass	pass	pass	pass	pass	pass

TABLE 3

	XIV	XV	XVI	XVII	XVIII
Component					
Oligomer C	50	60	50	50	50
Isobornylacrylate	14	14	20	24	
tricyclodecanedimethanol diacrylate	30		24	20	20
hexanedioldiacrylate		20			24
DAROCURE 1173 (Ciba-Geigy)	3.0	3.0	3.3	3.0	3.0
EBECRYL 170 (Radcure)	3.0	3.0	3.3	3.0	3.0
Viscosity (mPa.s)	1910	785	700	1240	1090
Dissipation factor at 25°C, 60Hz	0.020	0.029	0.019	0.024	~0.02
Dissipation factor at 150°C, 60Hz	0.083	0.032	0.060	0.082	~0.08
stability under 150°C oil immersion	pass	pass	slight defect	pass	pass
180° bend test	pass	pass	pass	pass	pass

TABLE 4

Temperature (°C)	IX	X	XI	XIII
25	0.029	0.019	0.0244	0.0246
36	0.031	0.022	0.0269	0.0277
45	0.033	0.026	0.0304	0.0305
55		0.026	0.0341	0.0311
65	0.024	0.027	0.0328	0.0297
75	0.017	0.023	0.029	0.0266
85	0.011	0.019	0.0254	0.0229
95	0.0065	0.014	0.0225	0.0208
105	0.006	0.011	0.0198	0.0197
115	0.007	0.011	0.0211	0.02
125	0.01	0.012	0.0258	0.0226
135	0.014	0.017	0.0358	0.0276
145	0.024	0.031	0.0566	0.0399
155	0.041	0.063	0.107	0.0663
165	0.07	0.14	0.168	0.114
175	0.126	0.31	0.29	0.233
185	0.27	0.78	0.579	0.401

While the invention has been described in detail and with reference to specific embodiments thereof, it will be apparent to those of ordinary skill in the art that various changes and modifications can be made therein without departing from the spirit and scope thereof.

What is claimed is:

1. A metal conductor having a cured coating of about 2.5 μm to about 500 μm thickness which coating has a dielectric dissipation factor (60Hz, 24°C) of lower than about 0.05 and is a radiation-cured coating formulated from a coating composition comprising:
 - a) an acrylate functional urethane oligomer having a hydrocarbon backbone;
 - b) at least one mono- or polyfunctional diluent; and optionally
 - c) a photoinitiator.
2. A metal conductor according to claim 1, wherein the metal is iron, copper, aluminum or silver.
3. A metal conductor according to claim 2, wherein the metal is aluminum, copper or silver.
4. A metal conductor according to claim 1, wherein the cured coating has an elongation at 25°C of greater than about 50% as a 25 μm thin cured coating.
5. A metal conductor according to claim 4, wherein the cured coating has at least one T_g of below 20°C as measured by the peak of the $\tan \delta$ curve in a

DMA analysis at 1 Hz.

6. A metal conductor according to claim 1, wherein the cured coating has a dissipation factor at 60 Hz at 150°C of lower than about 0.2.
7. A metal conductor according to claim 1, wherein the cured coating is formulated from the coating composition which further comprises an adhesion promotor.
8. A metal conductor according to claim 1, wherein the cured coating is a radiation-cured coating formulated from components comprising:
 - a) about 20 wt.% to about 80 wt.% of an acrylate functional urethane oligomer having a hydrocarbon backbone;
 - b) about 20 wt.% to about 80% of one or more mono- or polyacrylate functional monomers and optionally;
 - c) about 1 wt.% to about 10 wt.% of one or more light sensitive radical generating compounds; and optionally
 - d) about 0.2-5 wt.% of an adhesion promotor.
9. A metal conductor according to claim 1, wherein the cured coating is a coating cured by irradiation with light with a wavelength between about 200 μm to about 700 μm , and in the

photoinitiator is present in about 1 wt.% to about 10 wt.% of the coating composition.

10. A metal conductor according to claim 1, wherein the cured coating is a coating which is formulated from components comprising:
- a) about 30 wt.% to about 65 wt.% of an acrylate functional urethane oligomer having a hydrocarbon backbone;
 - b) about 20 wt.% to about 70 wt.% of at least two acrylate functional diluents, one of said acrylate functional diluents being a monoacrylate and one of said acrylate functional diluents being a polyacrylate functional compound;
 - c) about 1 wt.% to about 10 wt.% of one or more light sensitive radical generating compounds; and
 - d) about 0.2 wt.% to about 5 wt.% of an acid functional adhesion promotor.
11. A metal conductor according to claim 6, wherein the adhesion promotor is an acid functional compound.
12. A metal conductor to claim 1, wherein the cured coating further comprises about 0.2 wt.% to about 5 wt.% of a pigment or a dye.

13. A metal conductor according to claim 1, wherein the cured coating has a thickness of about 10 μm to about 100 μm .
14. A metal conductor according to claim 1, wherein the cured coating has a dielectric constant lower than about 5.
15. A metal conductor according to claim 1, wherein the cured coating has a dielectric dissipation factor lower than about 0.05 (60Hz, 24°C) after hot oil exposure (150°C).
16. A metal conductor according to claim 1, wherein the coating is formulated from components consisting essentially of:
- a) about 30 wt.% to about 65 wt.% of an acrylate functional urethane oligomer having a hydrocarbon backbone;
 - b1) about 10 wt.% to about 50 wt.% of a mono-acrylate functional diluent;
 - b2) about 5 wt.% to about 40 wt.% of a poly-acrylate functional diluent;
 - c) about 2 wt.% to about 7 wt.% of at least one photoinitiator;
 - d) about 0.2 wt.% to about 4 wt.% of adhesion promotor; and

e) about 0.2 wt.% to about 2 wt.% of a pigment.

17. A metal conductor according to claim 1, wherein said hydrocarbon backbone is fully saturated.

18. A radiation-curable coating composition comprising:

- a) an acrylate functional urethane oligomer having a hydrocarbon backbone;
- b) one or more mono- or polyfunctional diluents; and optionally
- c) one or more light sensitive radical generating compounds;

which coating when cured with radiation has a dielectric dissipation factor at 60 Hz at 24°C of lower than about 0.05, a dissipation factor at 60 Hz at 150°C of lower than about 0.2, and an elongation at 25°C of a 25 μ m thin cured coating of at least about 50%.

19. A radiation-curable coating composition according to claim 18, wherein said hydrocarbon backbone is fully saturated.

20. A radiation-curable coating according to claim 16, wherein the urethane oligomer is the reaction product of a hydrocarbon polyol, a polyisocyanate and an hydroxyfunctional endcapping monomer.

21. A radiation-curable coating composition according to claim 20, wherein said polyisocyanate is an aliphatic polyisocyanate.
22. A metal conductor according to claim 8, wherein said one or more monomers is a mono- or polyfunctional alkylacrylate or methacrylate based monomer.
23. A method of making a metal conductor with a cured coating of about 2.5 μm to about 500 μm thickness, which cured coating has a dielectric dissipation factor (60Hz, 24°C) of lower than about 0.05 comprising the steps of:
- a) providing a metal conductor;
 - b) coating said metal conductor with a radiation-curable coating composition which comprises;
 - i) an acrylate functional urethane oligomer having a hydrocarbon backbone;
 - ii) at least one mono- or polyfunctional diluent; and optionally
 - iii) a photoinitiator.

ABSTRACT OF THE DISCLOSURE

A metal conductor is provided with a cured coating of about 2.5 μm to about 500 μm thickness, which cured coating has a dielectric dissipation factor (60Hz, 24°C) of lower than about 0.05 and is a radiation-cured coating formulated from components comprising: a) an acrylate functional urethane oligomer having a hydrocarbon backbone; b) one or more mono- or polyfunctional diluents; and optionally, c) one or more light sensitive radical generating compounds. The invention further provides a radiation-curable coating composition for coating a metal conductor. Also disclosed is a method of making a metal conductor provided with a cured coating of about 2.5 μm to about 500 μm thickness, which cured coating has a dielectric dissipation factor (60Hz, 24°C) of lower than about 0.05.

46080T* 480T9680

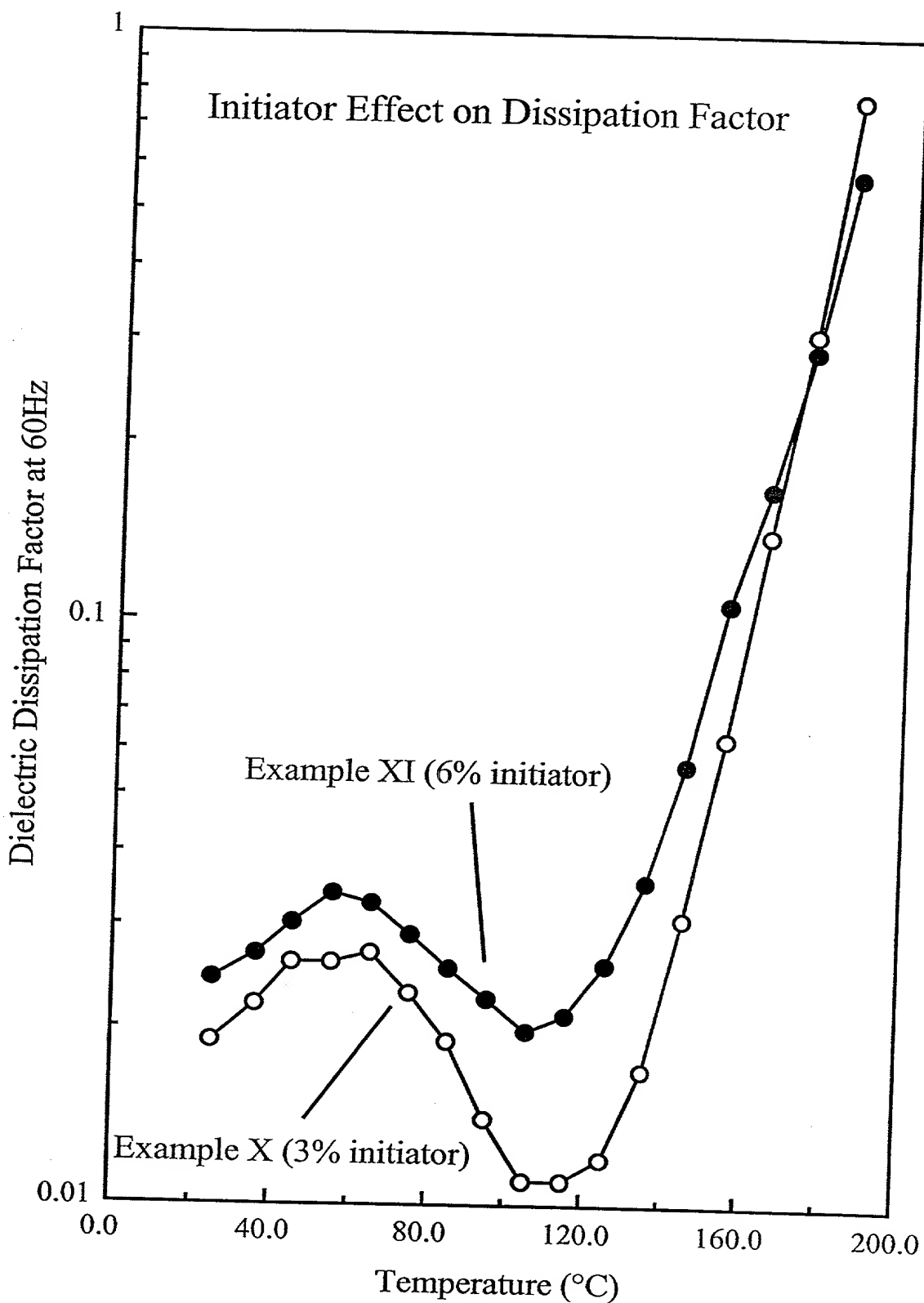


FIG 1

45030T, 430T, 5630

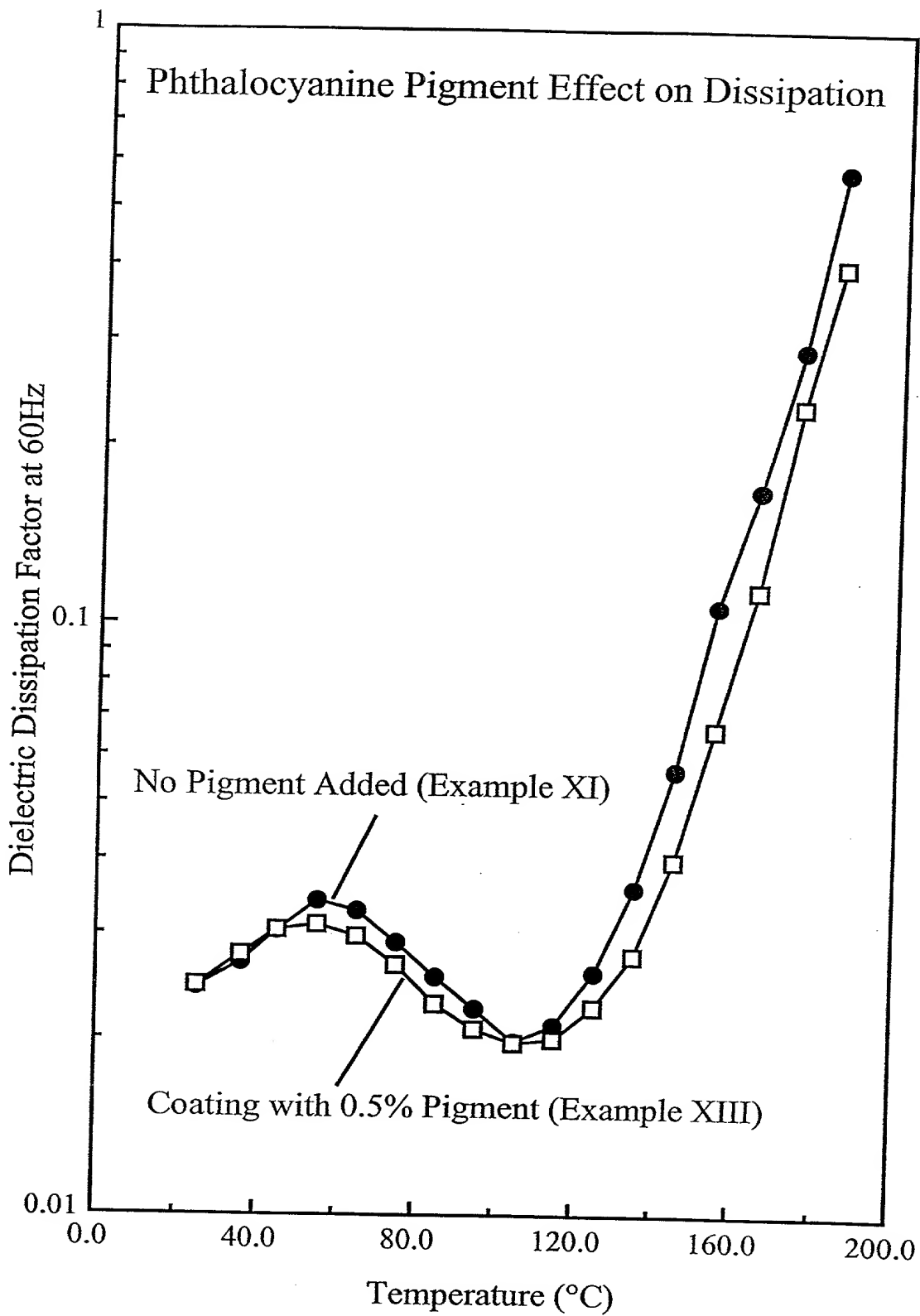
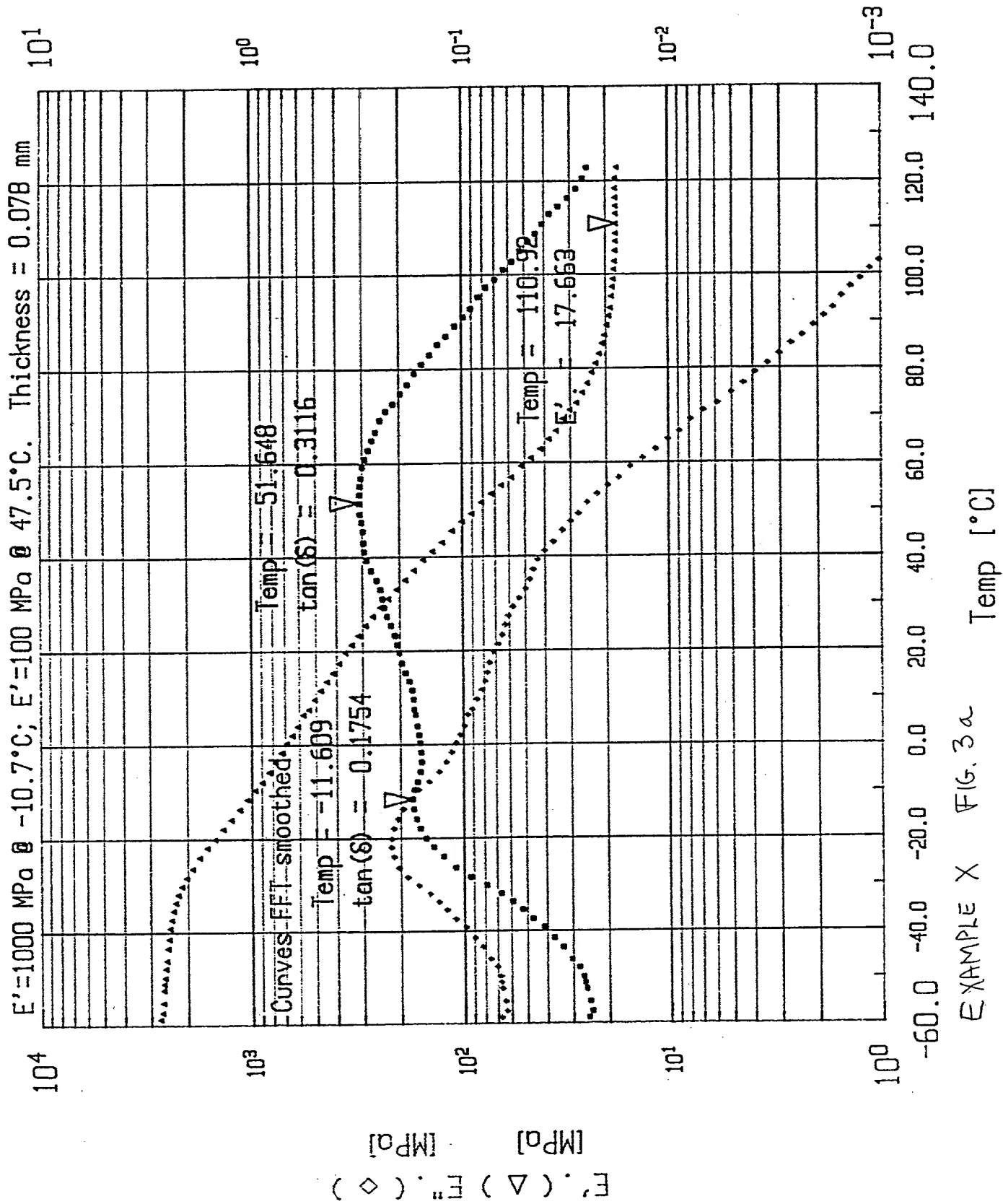
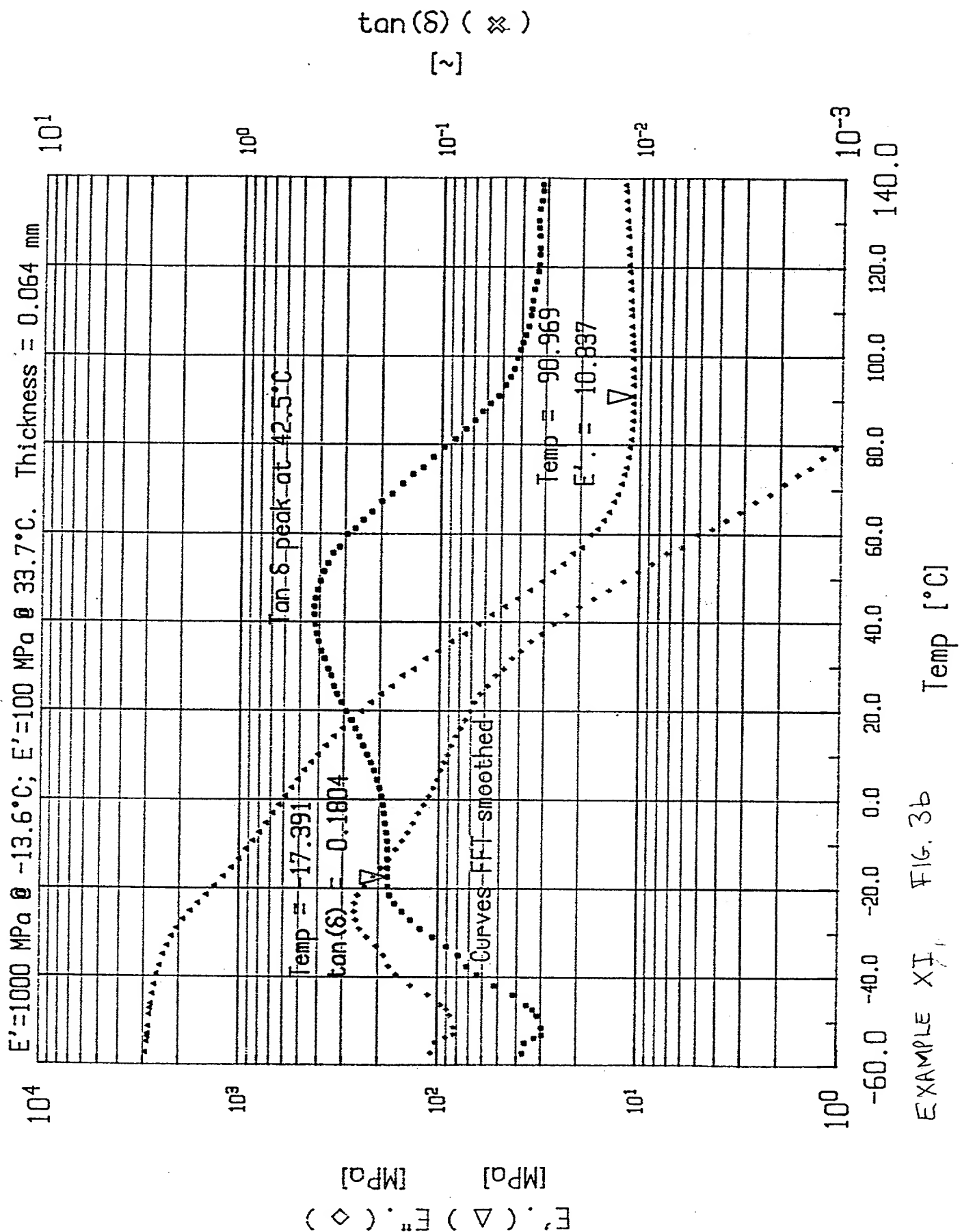


FIG. 2



EXAMPLE X FIG. 3a

60EOT 480T9680



EXAMPLE XI, FIG. 3b

Dielectric Coating on Aluminum Pannel (Example XI)

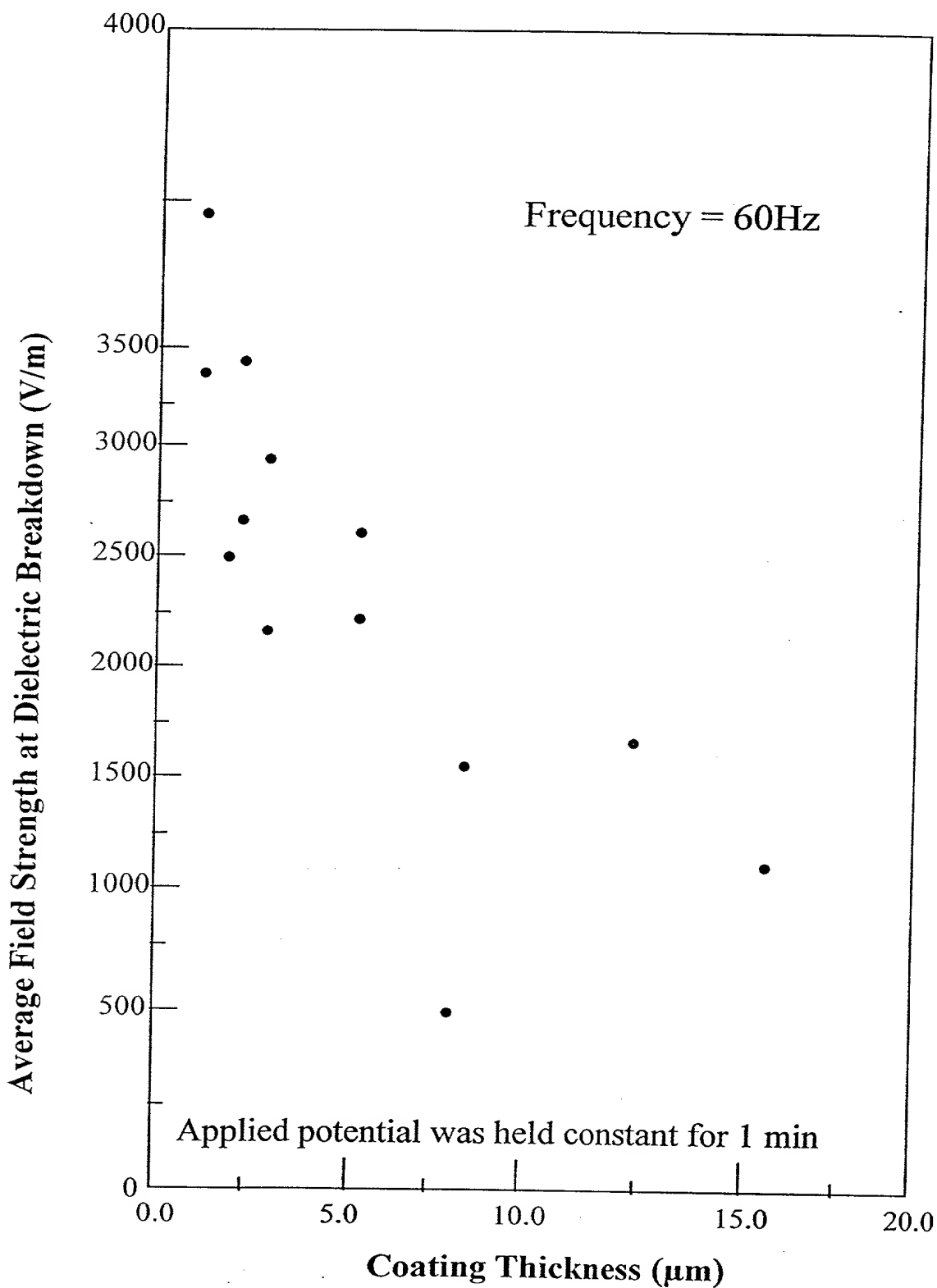


FIG. 4a

Dielectric Coating on Aluminum Pannel (Example XI)

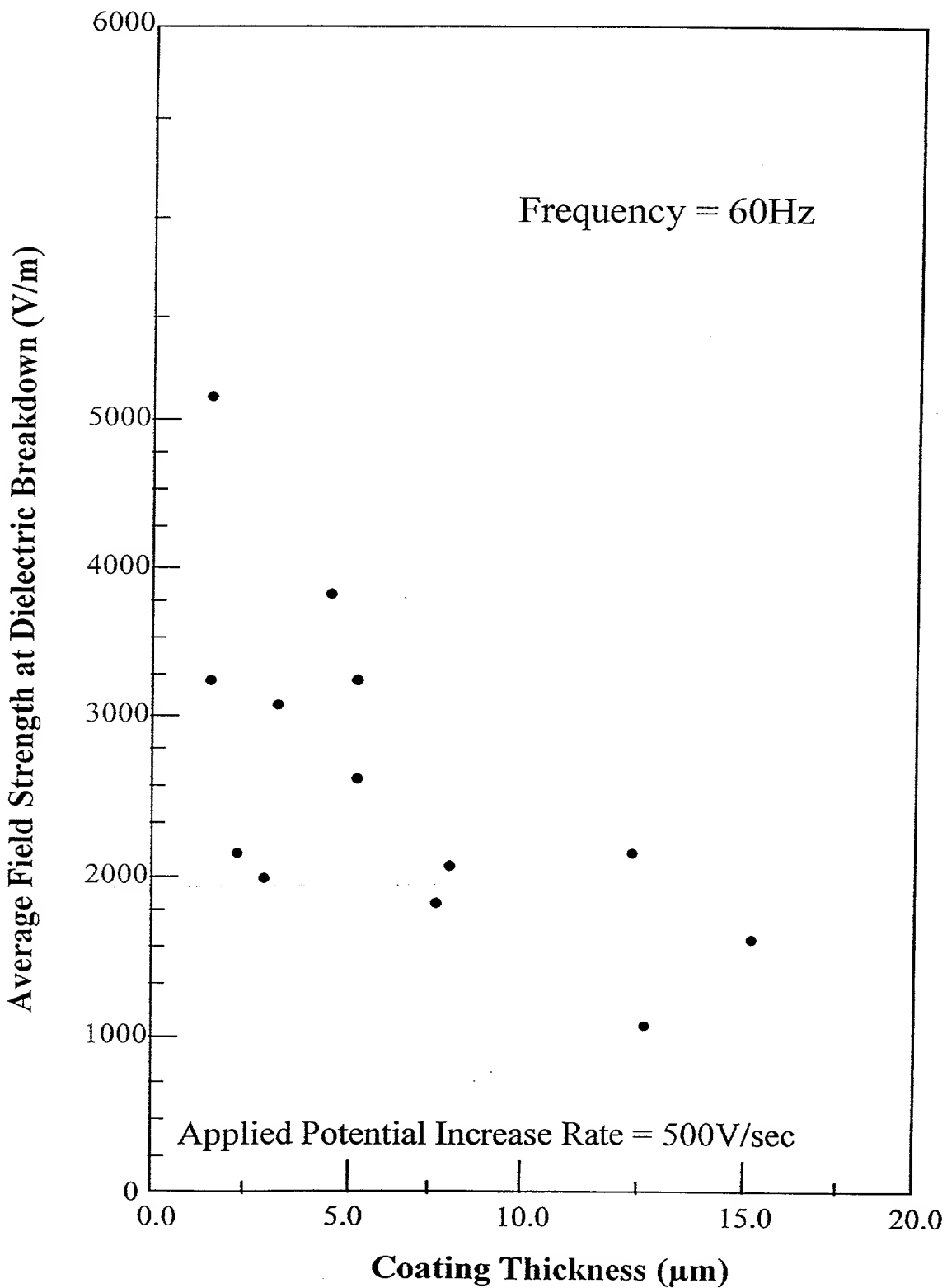


FIG. 4b